AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1. (currently amended): A pharmaceutical liquid composition comprising <u>5-methyl-1-phenyl-</u>2-(1H)-pyridone (Pirfenidone)a pyridone derivative represented by the following formula (I):

$$\frac{}{} R^2 - N$$

wherein R¹ is an alkyl group optionally having a substituent selected from the group consisting of a C₁₋₆ lower alkyl group optionally substituted at any of the 3-, 4- or 5-position with a halogen atom, a carboxyl group, an alkoxycarbonyl group, and an amino group and R² is a phenyl group optionally having a substituent selected from the group consisting of a C₁₋₆ lower alkyl group, a halogen atom, a carboxyl group, an alkoxycarbonyl group or an amino group, or a pharmaceutically acceptable salt thereof, and a solvent capable of dissolving said 5-methyl-1-phenyl-2-(1H)-pyridone (Pirfenidone)pyridone derivative in a concentration of 10% to about 25% by weight, wherein the solvent is diethylene glycol monoethyl ether.

- 2. (canceled).
- 3. (canceled).

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4. (currently amended): A pharmaceutical liquid composition according to Claim $3\underline{1}$, wherein

the diethylene glycol monoethyl ether has a purity of 99% or higher.

5. (previously presented): A pharmaceutical liquid composition according to Claim 1, further

comprising a concentrating agent.

6. (previously presented): A pharmaceutical liquid composition according to Claim 1, further

containing an antioxidant.

7. (original): A pharmaceutical liquid composition according to Claim 6, wherein the

antioxidant is an α -tocopherol.

8. (previously presented): A pharmaceutical liquid composition according to Claim 1, in the

form of an oral, percutaneous, nasal or vaginal preparation or in the form of a spray, patch,

inhalant, injection or intravenous drip.

9. (previously presented): A pharmaceutical liquid composition according to Claim 1, having

the following components:

<u>Ingredients</u> % by weight

Pirfenidone 10-25

Diethylene glycol

monoethyl ether 70-80

Ethanol (95%) 0-10

Polyvinyl pyrrolidone or

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hydroxypropyl cellulose 0-3

Sodium metabisulfite 0.02-2

Methyl or propyl

paraben 0-0.5

Purified water 0-25.

10. (previously presented): A pharmaceutical liquid composition according to Claim 1, having the following components:

<u>Ingredients</u>	% by weight
Pirfenidone	10-25
Diethylene glycol	
monoethyl ether	75-80
Purified water	<u>0-10</u> .

11. (previously presented): A pharmaceutical liquid composition according to Claim 18, having the following components:

<u>Ingredients</u>	% by weight
Pirfenidone	10-25
Diethylene glycol	
monoethyl ether	75-80
α-Tocopherol	0.1-0.5
Hydroxypropyl cellu	lose 0-3
Purified water	0-10.

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12. (new): The pharmaceutical composition according to claim 1, wherein the composition has good stability.

13. (new): The pharmaceutical composition according to claim 1, wherein the composition does not cause skin irritation.